

**Supplementary Table 2: Investigations and treatment of vitamin D deficiency**<sup>2,3,5,24,25</sup>

<b>Vitamin D deficiency in congenital ichthyosis</b>		
<ul style="list-style-type: none"> <li>- Vitamin D, a fat-soluble vitamin, has its first prohormone being synthesised in skin after exposure to ultraviolet B (UVB) light.</li> <li>- Initial correction and maintenance of vitamin D and calcium does not require urinary calcium:creatinine ratio. In addition to improvement in bone health, this takes around 4–10 weeks depending on the severity; improvement in cutaneous disease is also documented, around ten days as noticed by Sethuraman G <i>et al.</i></li> <li>- In a recent study conducted by Mahajan <i>et al.</i>, although there was a statistically significant difference in mean vitamin D levels between the ichthyosis phenotypes, there was no correlation between the serum vitamin D levels and ichthyosis severity scores, thus emphasising the necessity of screening for Vitamin D deficiency in all congenital ichthyosis patients, including those with milder phenotypes.</li> </ul>		
<b>Investigations</b>		
<b>Hormonal</b>	<ul style="list-style-type: none"> <li>- Serum 25-hydroxyvitamin D (25OH-D)</li> <li>- Serum parathyroid hormone (PTH)</li> </ul>	
<b>Biochemical</b>	<ul style="list-style-type: none"> <li>- Alkaline phosphatase (ALP)</li> <li>- Serum calcium (Ca<sup>2+</sup>)</li> <li>- Serum phosphate</li> <li>- Urinary calcium-creatinine ratio</li> </ul>	
<b>Radiological</b>	<ul style="list-style-type: none"> <li>- X-ray distal wrist and knees (sites of rapid bone growth)</li> <li>- Other sites (costochondral junction, skull)</li> <li>- Dual energy X-ray absorptiometry scan (DEXA scan)</li> </ul>	
<b>Treatment</b>		
	<b>Vitamin D &lt;20 ng/mL WITH clinical rickets</b>	<b>Vitamin D &lt;20 ng/mL WITHOUT clinical rickets</b>
<b>Intensive phase</b>	Cholecalciferol 60,000 IU weekly for 10 weeks + elemental Ca <sup>2+</sup> 50–75 mg/kg/day Review in 12 weeks with repeat X-rays and labs <ul style="list-style-type: none"> <li>- Healed → maintenance phase</li> <li>- Not healed → continue till radiological healing seen</li> </ul>	Cholecalciferol 60,000 IU weekly for four weeks + elemental Ca <sup>2+</sup> 50–75 mg/kg/day
<b>Maintenance phase</b>	Cholecalciferol 400 IU (infants) – 600 IU (children >1 year) daily + elemental Ca <sup>2+</sup> 50–75 mg/kg/day to be given lifelong Review in three months with 25-OH-D and PTH	Cholecalciferol 400 IU (infants) – 600 IU (children >1 year) daily OR Cholecalciferol 60,000 IU monthly + elemental Ca <sup>2+</sup> 50–75 mg/kg/day to be given lifelong
<b>Follow-up</b>	Yearly bone and lab parameters	Yearly lab parameters
<p style="text-align: center;"><b>Stoss therapy for vitamin D deficient rickets</b></p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Particularly helpful for patients with poor compliance to therapy</li> <li>- Oral administration of 100,000–600,000 IU vitamin D daily (over a period of 1–5 days) followed by eight weeks of 400–1000 IU Vitamin D daily OR 50,000 IU Vitamin D weekly</li> </ul>		

*UVB, Ultraviolet B; 25OH-D, 25-hydroxycholecalciferol; PTH, Parathyroid hormone; ALP, Alkaline Phosphatase; Ca<sup>2+</sup>, Calcium; DEXA, Dual energy X-ray absorptiometry; IU, International units.*

#### **SUPPLEMENTARY FILES: Additional references**

24. Gupta P, Dabas A, Seth A, Bhatia VL, Khadgawat R, Kumar P, et al. Indian Academy of Pediatrics Revised (2021) Guidelines on Prevention and Treatment of Vitamin D Deficiency and Rickets. *Indian Pediatr* 2022;59(2):142-58.
25. Mahajan R, Bakshi S, Chatterjee D, De D, Saikia UN, Handa S. Clinico-epidemiologic profile of non-syndromic congenital ichthyosis - a retrospective chart review of 107 Patients. *Indian J Dermatol* 2024;69:113-8.